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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/823,012	04/13/2004	Micheal Patrick Dillon	R0130D-CON	4690
24372 ROCHE PALO	7590 02/12/2007 ALTO LLC		EXAM	INER
PATENT LAW	DEPT. M/S A2-250		STOCKTON, LA	AURA LYNNE
3431 HILLVIE PALO ALTO, O			ART UNIT	PAPER NUMBER
11.20.22.0,			1626	
SHORTENED STATUTORY	Y PERIOD OF RESPONSE	MAIL DATE	DELIVER'	Y MODE
3 MOI	NTHS	02/12/2007	PAP	ER

Please find below and/or attached an Office communication concerning this application or proceeding.

If NO period for reply is specified above, the maximum statutory period will apply and will expire 6 MONTHS from the mailing date of this communication.

	Application No.	Applicant(s)
Office Action Commons	10/823,012	DILLON ET AL.
Office Action Summary	Examiner	Art Unit
	Laura L. Stockton, Ph.D.	1626
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the c	orrespondence address
A SHORTENED STATUTORY PERIOD FOR REPLY THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.1: after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a reply - If NO period for reply is specified above, the maximum statutory period of the period for reply within the set or extended period for reply will, by statute any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	36(a). In no event, however, may a reply be tim y within the statutory minimum of thirty (30) days will apply and will expire SIX (6) MONTHS from to cause the application to become ABANDONE	nely filed s will be considered timely. the mailing date of this communication. D (35 U.S.C. § 133).
Status		
1) Responsive to communication(s) filed on 29 N	ovember 2006.	
2a)⊠ This action is FINAL . 2b)☐ This	action is non-final.	
3) Since this application is in condition for alloward closed in accordance with the practice under E	·	
Disposition of Claims		
4) ⊠ Claim(s) 48 and 51-62 is/are pending in the ap 4a) Of the above claim(s) 53 and 55-62 is/are v 5) □ Claim(s) is/are allowed. 6) ⊠ Claim(s) 48,51,52 and 54 is/are rejected. 7) □ Claim(s) is/are objected to. 8) □ Claim(s) are subject to restriction and/o	vithdrawn from consideration.	
Application Papers		N.
9) The specification is objected to by the Examine	ır.	
10) ☐ The drawing(s) filed on is/are: a) ☐ acc	epted or b) \square objected to by the $\mathfrak k$	Examiner.
Applicant may not request that any objection to the	• • • • • • • • • • • • • • • • • • • •	
Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Ex		
Priority under 35 U.S.C. § 119		
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the priority application from the International Bureau * See the attached detailed Office action for a list	s have been received. s have been received in Applicati rity documents have been receive u (PCT Rule 17.2(a)).	on No ed in this National Stage
Attachment(s)		
1) Notice of References Cited (PTO-892)	4) Interview Summary Paper No(s)/Mail Da	
 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08) Paper No(s)/Mail Date 12/11/2006. 		Patent Application (PTO-152)

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DETAILED ACTION

Claims 48 and 51-62 are pending in the application.

Election/Restrictions

Applicant's election without traverse of Group I, and the species of Example 3 in step 4 on page 49 (reproduced below), in the reply filed on December 16, 2004 was acknowledged in a previous Office Action.

Step 4

The requirement was deemed proper and made FINAL in a previous Office Action.

Claims 53 and 55-62 are withdrawn from further consideration pursuant to 37 CFR 1.142(b) as being

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drawn to nonelected inventions. Election was made without traverse in the reply filed on December 16, 2004.

Information Disclosure Statement

The Examiner has considered the Information Disclosure Statement filed on December 11, 2006.

Rejections made in the previous Office Action that do not appear below have been overcome. Therefore, arguments pertaining to these objections will not be addressed.

Terminal Disclaimer

The terminal disclaimer filed on May 10, 2006 disclaiming the terminal portion of any patent granted on this application which would extend beyond the

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expiration date of U.S. Patent 6,756,395 has been reviewed and is accepted. The terminal disclaimer has been recorded.

The Declaration under 37 CFR 1.132 filed

May 10, 2006 by Counde O-Yang has been re-assessed

since Applicant has provided further explanation of the showing; has submitted expanded forms of Figure 1 and

Figure 2; and has amended the claims.

Response to Amendment

The Declaration under 37 CFR 1.132 filed

May 10, 2006 by Counde O-Yang is insufficient to

overcome the rejection of claims 48, 51, 52 and 54

based upon 35 USC 103 as set forth in the last Office

action because the showing is not commensurate in scope

with the currently amended claims. *In re Greenfield*,

197 U.S.P.Q. 227 (1978) and *In re Lindner*, 173 U.S.P.Q.

356 (1972). Also see M.P.E.P. 716.02(d).

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Applicant's showing is not commensurate in scope because Applicant has only compared one of the compounds of the instant claimed invention with only one compound of Cournoyer et al. The comparison of only one compound is insufficient to show a true trend that the para compounds have unexpected and beneficial properties over the meta compounds. By comparing the compounds found in Table 1 of the instant specification starting at page 19 (reproduced below) with the compounds of Cournoyer et al. (relevant columns reproduced below), the following table illustrates some of the other compounds which should also have been compared to persuasively show that the instant claimed compounds have unexpected, beneficial and unobvious results over the compounds of Cournoyer et al. in treating urinary incontinence while not increasing blood pressure. See, for example, the table below.

Applicants' Compounds	Cournoyer et al. Compounds				
Compound 2▶	The Compound in column 40, lines 10-12				

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Compound 3	 The	Compound	in	column	39,	lines	16-18
Compound 4	 The	Compound	in	column	39,	lines	38-40
Compound 5	 The	Compound	in	column	39,	lines	32-34
Compound 29	 						
Compound 40	 The	Compound	in	column	39,	lines	46-48

Instant Compounds found on pages 19 and 21-24, respectively, of the instant specification that are embraced by independent claim 48 follow.

2	N-[4-(4,5-Dihydro-1H-imidazol-2-ylmethyl)-2-methoxy-phenyl]-methanesulfonamide;	2	OSS N
3	N-[4-(4,5-Dihydro-1H-imidazol-2-ylmethyl)-2-methyl-phenyl]-methanesulfonamide	2	H ₃ C N N
4	N-[2-Chloro-4-(4,5-dihydro-1H-imidazol-2-ylmethyl)-phenyl]-methanesulfonamide	3	OSS N
5	N-[4-(4,5-Dihydro-1H-imidazol-2-ylmethyl)-2-hydroxy-phenyl]-methanesulfonamide	2	H ₃ C S N N N
29	N-[2-Ethoxy-4-(4,5-dihydro-1H-imidazol-2-ylmethyl)-phenyl]-methanesulfonamide	4	H ₃ C O N N

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L			п п
33	N-[2-Bromo-4-(4,5-dihydro-1H-imidazol-2-ylmethyl)-phenyl]-methanesulfonamide	4	Br O NH
40	N-[4-(4,5-Dihydro-1H-imidazol-2-ylmethyl)-2-fluoro-phenyl]-methanesulfonamide	4	H ₃ C N N N

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Compounds of Cournoyer et al. found in Columns 39 and 40.

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39

$$R^{23}$$
 R^{24}
 $R^{22}SO_2N$
 R^{24}
 R^{25}
 R^{25}

wherein \mathbf{Y}^{c} is 2-imidazoline, and

			<u> </u>		
R ²²	R ²³	R ²⁴	R ²⁵	R ²⁶	R ²⁷
CH ₃	H	CH,	H	н	H
(N-[3-(4.5)	-dihydro-11	H-imidazol-2	-yimethyl)-2	2-methyl-ph	enyl]-
methanesu	ılfonamide)	•			
CH ₃	H	H	H	H	CH₃
(N-[5-(4,5)]	-dihydro-11	H-imidazol-2	-ylmethyl)-2	2-methyl-ph	enyl]-
methanesu	ılfonamide)	•			
CH ₃	H	H	H	Cl	CH ₃
N-[3-chlor	ro-5-(4,5-di	hydro-1H-im	ridazol-2-ylr	nethyl)-2-m	ethyl-phenyl]-
	lfonamide		•		
CH ₂	H	H	H	Br	CH₃
(N-13-bron	mo-5-(4.5-d	lihydro-1H-i	midazol-2-yl	methyl)-2-	nethyl-phenyl]-
	lfonamide)		•	- -	· ·
CH ₃	H	н	H	H	OCH_3
N-15-14.5		H-imidazol-2		2-methoxy-	
	lfonamide)		· , , -, -	·- · · · · · · · · · · · · · · ·	
CH ₃	H	H	Ħ	H	H
(NI_[3_(/ 5		H-imidazol-2			
	lfonamide)		·		
	H H	, H	н	н	ОН
CH ₃	ia Saibudyo 11	H-imidazol-2			
			-yimicemyi)**	L Lyuluny-1	,.,
	ilfonamide)	H	F	н	H
CH ₃	H				
		H-imidazol-2	-yimethyi)	+-11GO1O-pii	311313-
	ilfonamide)		77	**	H
CH ₃	CH₃	CH,	H	H	
		H-imidazol-2	!-yimethyi)-	z-metnyi-pi	ienyij- wi v:-
methyl-me	ethanesulfo				
CH_3	H	Cl	H	H	H
(N-[2-chlo	oro-3-(4, 5 -d	lihydro-1H-i:	midazol-2-yl	methyl)-ph	enyl]-
methanest	ılfonamide))			
CH ₃	H	C _o H _s	H	H	H
(N-16-(4.5)	-dihydro-1	H-imidazol-2	2-ylmethyl)-i	biphenyl-2-	y1]-
	lfonamide)				
CH ₃	H	CH ₃	CH_3	H	H
(N-[3-(4.5	-dihydro-1	H-imidazol-2	2-ylmethyl)-2	2,4-dimethy	d-phenyl]-
	lfonamide)			_	
CH ₂	H	H	H	H	F
(N-15-(4.5		H-imidazol-2	2-ylmethyl)-:	2-fluoro-ph	enyl]-
	lfonamide		,	•	- -
CH ₃	H	CH,	H	H	CH3
(NI-13-(4 5	(-dibydro-1	H-imidazol-2			
	alfonamide			_,,	- 1
	H H	CH ₃	H	CH_3	Ħ
CH ₃	Fl : diby:dec 4	CH₃ H-imidazol-2		2 6-dimeths	
			- Finterny 1)-	2,0-umicu)	- p
	ulfonamide)		L H	н	H
CH ₃	. н	CH=CH			**
		4-ylmethyl)-2	c-metnyi-ph	enyı J-	
	ulfonamide				T-T
CH ₂	H	C_2H_{ϵ}	H	H	H

40 -continued

	R ²²	R ²³	R ²⁴	R ²⁵	R26	R ²⁷
	methanesui	ionamid)				
5	CH,	H	\mathbf{H}	H	OCH ₃	H
	(N-[3-(4,5-c		I-imidazol-2-	ylmethyl)-5	-methoxy-pl	henyi]-
	CH ₂ CH ₂	H	H	H	H	CH ₃
	(ethanesulfo	_	5-(4,5-dihydro	-1H-imida:	zel-2-ylmeth	yl)-2-methyl-
)	CH.	H	OCH ₂	H	H	H
	(N-[3-4,5-d methanesuli		imidazel-2-yl	methyl)-2-	methoxy-phe	nyl]-

Response to Arguments

Applicant's arguments filed November 29, 2006 have been considered. Applicant argues that the genus of the instant claimed compounds has been reduced to cover seven specific compounds since additional comparison data was not available. Applicant argues that the showing of the Rule 1.132 Declaration is now commensurate in scope with the currently amended claims. In response, the showing in the Declaration is insufficient for reasons stated above.

Double Patenting

The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the "right to exclude" granted by a patent and to prevent possible harassment by multiple assignees. See In re Goodman, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); In re Longi, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); In re Van Ornum, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); In re Vogel, 422 F.2d 438, 164 USPQ 619 (CCPA 1970); and, In re Thorington, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1.321(c) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent is shown to be commonly owned with this application. See 37 CFR 1.130(b).

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

Claims 48, 51, 52 and 54 are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claims 1-4, 8, 10, 12-14, 18, 20, 27-30, 32, 44-47, 56-58, 60 and 64 of U.S. Patent No. 5,952,362 (Cournoyer et al.). Although the conflicting claims are not identical, they are not

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patentably distinct from each other because the instant claimed compounds are positional isomers of the compounds claimed in U.S. Patent No. 5,952,362.

In U.S. Patent No. 5,952,362, see claim 1 (columns 124-125) and especially claim 27 (column 127) and claim 44 (column 128). The sulfonamide group in the compounds found in U.S. Patent No. 5,952,362 is attached to the phenyl ring meta to the imidazolin-2yl-methyl group instead of para to the imidazolin-2-ylmethyl group as instantly claimed (i.e., a positional Nothing unobvious is seen in substituting the known claimed isomer for the structurally similar isomer, as claimed in U.S. Patent No. 5,952,362 since such structurally related compounds suggest one another and would be expected to share common properties absent a showing of unexpected results. In re Norris, 84 USPQ 458 (1950).

One skilled in the art would thus be motivated to prepare positional isomers of the compounds claimed in

U.S. Patent No. 5,952,362, to arrive at the instant claimed compounds with the expectation of obtaining additional beneficial products which would be useful in treating, for example, urinary incontinence. The instant claimed invention would have been suggested and therefore, obvious to one skilled in the art.

Response to Arguments

Applicant's arguments filed November 29, 2006 have been considered. Applicant argues that Declaration under 37 CFR 1.132 filed May 10, 2006 by Counde O-Yang demonstrates unexpected better ability of the claimed compounds to increase IUP without causing an undesirable increase in blood pressure compared to the compounds of Cournoyer et al. In response, the Declaration shows that one of the instant claimed compounds demonstrates unexpected better ability to increase IUP without causing an undesirable increase in blood pressure compared to the compounds of Cournoyer

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et al. However, the Declaration under 37 CFR 1.132 filed May 10, 2006 by Counde O-Yang was still considered insufficient for reasons stated above.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

Claims 48, 51, 52 and 54 are rejected under 35 U.S.C. 103(a) as being obvious over Cournoyer et al. {U.S. Pat. 5,952,362} for reasons set forth below.

The applied reference has a common inventor (i.e., Counde O'Yang) with the instant application. Based

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upon the earlier effective U.S. filing date of the reference, it constitutes prior art only under 35 U.S.C. 102(e). This rejection under 35 U.S.C. 103(a) might be overcome by: (1) a showing under 37 CFR 1.132 that any invention disclosed but not claimed in the reference was derived from the inventor of this application and is thus not an invention "by another"; (2) a showing of a date of invention for the claimed subject matter of the application which corresponds to subject matter disclosed but not claimed in the reference, prior to the effective U.S. filing date of the reference under 37 CFR 1.131; or (3) an oath or declaration under 37 CFR 1.130 stating that the application and reference are currently owned by the same party and that the inventor named in the application is the prior inventor under 35 U.S.C. 104, together with a terminal disclaimer in accordance with 37 CFR 1.321(c). This rejection might also be overcome by showing that the reference is disqualified under 35

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U.S.C. 103(c) as prior art in a rejection under 35
U.S.C. 103(a). See MPEP § 706.02(l)(1) and §
706.02(l)(2).

Claims 48, 51, 52 and 54 are rejected under 35 U.S.C. 103(a) as being unpatentable over Cournoyer et al. {U.S. Pat. 5,952,362}.

Determination of the scope and content of the prior art (MPEP §2141.01)

Applicant claims imidazolin-2-yl-methylphenyl compounds. Cournoyer et al. teach imidazolin-2-yl-methylphenyl compounds that are structurally similar to the instant claimed compounds. See in Cournoyer et al., for example, formula 1 in columns 7 and 8 and especially the first compound listed in the table in column 39.

Ascertainment of the difference between the prior art and the claims (MPEP §2141.02)

The difference between the compounds of Cournoyer et al. and the compounds instantly claimed is that the sulfonamide group is attached to the phenyl ring meta to the imidazolin-2-yl-methyl group instead of para to the imidazolin-2-yl-methyl group as instantly claimed (i.e., a positional isomer).

Finding of prima facie obviousness--rational and motivation (MPEP §2142-2413)

Nothing unobvious is seen in substituting the known claimed isomer for the structurally similar isomer, as taught by Cournoyer et al., since such structurally related compounds suggest one another and would be expected to share common properties absent a showing of unexpected results. *In re Norris*, 84 USPQ 458 (1950).

One skilled in the art would thus be motivated to prepare positional isomers of the compounds taught by Cournoyer et al. to arrive at the instant claimed compounds with the expectation of obtaining additional beneficial products which would be useful in treating,

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for example, urinary incontinence. The instant claimed invention would have been suggested and therefore, obvious to one skilled in the art. A strong case of prima facie obviousness has been established.

Response to Arguments

Applicant's arguments filed November 29, 2006 have been considered. Applicant argues that Declaration under 37 CFR 1.132 demonstrates unexpected better ability of the claimed compounds to increase IUP without causing an undesirable increase in blood pressure compared to the compounds of Cournoyer et al. In response, the Declaration shows that one of the instant claimed compounds demonstrates unexpected better ability to increase IUP without causing an undesirable increase in blood pressure compared to the compounds of Cournoyer et al. However, the Declaration under 37 CFR 1.132 filed May 10, 2006 by Counde O-Yang

was still considered insufficient for reasons stated above.

Conclusion

Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will

be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

This application contains claims 53 and 55-62 drawn to an invention nonelected without traverse in the reply filed December 16, 2004. A complete reply to the final rejection must include cancellation of nonelected claims or other appropriate action (37 CFR 1.144) See MPEP § 821.01.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Laura L. Stockton whose telephone number is (571) 272-0710. The examiner can normally be reached on Monday-Friday from 6:15 am to 2:45 pm. If the examiner is out of the Office, the examiner's supervisor, Joseph McKane, can be reached on (571) 272-0699.

Information regarding the status of an application may be obtained from the Patent Application Information

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The Official fax phone number for the organization where this application or proceeding is assigned is (571) 273-8300.

Laura L. Stockton, Ph.D.

Patent Examiner

Art Unit 1626, Group 1620

Technology Center 1600

February 5, 2007